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(54) **Aromatic polycarbonate composition.**

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Description

Injection molding provides a convenient way for preparing various articles from thermoplastic resins, particularly objects of a relatively intricate nature. In order to injection mold articles in an economic manner, it is important to minimize the adhesion of the molded part to the mold surface through the use of a release agent. It is additionally advantageous that the release agent be incorporated in the material to be demolded. However, such an internal agent must be compatible with the resin, that is, it should not lead to any significant loss of the other physical properties such as transparency and melt stability as measured by the usual characteristics of the resin under normal conditions and heat treatments.

Of the thermoplastic resins which find a mold release agent useful from time to time are polycarbonate, copolyestercarbonate and blends of these resins with aliphatic polyester resins. As a result of the generally high temperature which these thermoplastics are subjected to during molding and the inherent sensitivity of these resins to chemical attack and degradation by additives which are often compatible with many other thermal plastics, it is necessary to select mold release agents with high temperature stability and compatibility with polycarbonates. Mineral oils are one types of mold release agents as described e.g. in GB-A-2,077,742.

Mineral oils are naturally occurring complex mixtures of paraffinic and naphthalenic hydrocarbons that are obtained from petroleum.

We have found that synthetic mineral oils derived from the oligomerization of alpha olefins followed by hydrogenation to remove unsaturation provide a surprising benefit over normal paraffins and conventional mineral oils as mold release agents in various polymers including polycarbonate, copolyestercarbonate and blends of polycarbonate with other resins including cycloaliphatic or aliphatic-cycloaliphatic polyester resins without introducing significant detrimental effects on color, stability and other properties associated with these resins.

In accordance with the invention, there is a composition comprising an aromatic carbonate polymer in admixture with a mold release effective amount of a hydrogenated alpha-olefin oligomer fluid. Although a wide range of alpha olefins and mixtures of olefins can be used to prepare hydrogenated alpha-olefin oligomeric fluids, the preferred alpha olefin is 1-decene.

The aromatic carbonate polymers are prepared by well known techniques available in the art, see for example U.S.-A-4,444,935, column 2, line 26 to column 3, line 61. The typical polycarbonates can be employed in the invention, for example, homopolycarbonates such as bisphenol-A polycarbonates and copolycarbonates wherein two or more dihydric phenols are employed. Further aromatic carbonate polymers include the copolyestercarbonates such as those disclosed in U.S.-A-3,169,121 and 4,465,820.

The hydrogenated alpha-olefin oligomer fluids useful in this invention are derived from the oligomerization of 1-olefins through a catalytic reaction followed by hydrogenation to remove unsaturation. Examples of such catalysis include: cationic catalysis, see J.A. Breenan, Ind. Eng. Chem. Prod. Res. Div. (1980) 19, 2; Ziegler catalysis, see D.H. Antonsen et al, Ind. Eng. Chem. Prod. Res. Div. (1963), 2, 224; or metathesis catalysis, see W.T. Nelson et al, Ind. Eng. Chem. Prod. Res. Div., (1983), 22, 178 and references therein. Methods of preparation are also disclosed in U.S.-A-4,282,392; 4,225,739; 4,319,065; 4,311,864; 4,334,113 and 4,409,415 and references therein.

The product of oligomerization/hydrogenation is a mineral oil with fluidity over a wide temperature range. The product generally is a mixture of oligomers of various degrees of polymerization, branching and isomerization depending on the nature of the catalyst and olefin used and any subsequent product fractionation such as by distillation. A product structure representative of this type of oligomerization is 9-methyl-11-n-octyl heneicosane. Normal alpha olefins of 6-12 carbon atoms are preferred for the preparation of synthetic fluids, with 1-decene being most common. The oligomers are generally separated into various mixtures of dimer, trimer, tetramer and pentamer to achieve varying degrees of volatility, viscosity and viscosity index. The preferred saturated alpha-olefin oligomer fluids for this invention are those with a kinematic viscosity of approximately 2-8 mm²/s (2-8 cSt) at 100°C. as determined by ASTM D445. For the purposes of this invention, the saturated alpha-olefin oligomer fluids can also be blended with other lubricants such as the synthetic and naturally occurring polyol esters.

An effective mold releasing amount of the material of the invention is employed in the thermoplastic composition. Any amount of material which reduces the amount of pressure needed to eject the article from the injection mold and obtain a substantially unblemished article in comparison to the pressure needed to eject the thermoplastic composition control is an effective mold releasing amount. In general, effective amounts of the material are from 0.01 to 1.0 weight percent, based on the quantity of thermoplastic resin present, preferably from 0.05 to about 0.5 weight percent. The material can be added to the resin in the normal manner that the other additives are added and coextruded. Depending upon the volatility of the

compound at common processing and extrusion temperatures, an excess may be initially employed in the composition.

Other common additives for thermoplastic resins may also be employed. For example with respect to aromatic polycarbonate, additives which are commonly known to stabilize the resin thermally such as a phosphite can be employed. Stabilizers such as epoxides may also be employed as well as agents which are flame retardants, drip inhibitors, ductility enhancers, antioxidants, solvent resistance enhancers, ultraviolet light stabilizers, colorants and various inert fillers. Impact modifiers may also be present in the composition.

Below are specific examples of the invention. The examples are intended to illustrate but not narrow the inventive concept. The saturated alpha-olefin oligomer fluids used in the following examples are those available from Emery Industries. These synthetic fluids are derived from 1-decene and are referred to as Emery 3002, Emery 3004 and Emery 3006. The approximate composition of these fluids, as determined by gas chromatography is shown below.

	DIMER	TRIMER	TETRAMER	PENTAMER	AVG.# CARBONS
3002	100%	---	---	---	20
3004	---	89	11	---	31
3006	---	29	62	9	38

Alternatively, the Synfluids manufactured by Gulf Oil Corporation could also be used. All percentages of the additives are in weight percent of the thermoplastic resin.

In the ensuing examples the aromatic polycarbonate is LEXAN® 140, a polymer produced by reacting bisphenol-A and phosgene and having an intrinsic viscosity at 25°C. in methylene chloride at 25°C. of 0.49-0.52 dl/grams. The copolyester carbonate is prepared in accordance with the methods of U.S.A-4,465,820 wherein bisphenol-A is the dihydric phenol and has 75 mole percent phthalate ester linkages of which 7 percent are terephthalate, the remainder isophthalate. Kodar® A150 is available from Eastman and is a polyester of trans 1,4-cyclohexane dimethanol and iso and terephthalic acid. The mold release agent was dry formulated into the resins at levels of 0.3 weight percent unless otherwise stated. Also present in the composition was 0.03 weight percent of a phosphite stabilizer.

Mold release effectiveness was determined in an injection molding machine with a 124.4 g (4 ounce) shot capacity. The part molded was a 7.62 cm x 7.62 cm (3" x 3") box with a 3.81 cm (1 1/2") wall height. It had ejector pins at four corners with two of the pins being attached to strain gauge transducers for measuring the part ejection pressure. The mold was designed with very little draft so that the part would tend to stick to the core unless mold release is used. To vary the amount of shrinkage on the core, the temperature of the stationary and moving mold halves could be varied. The ejector pin pressure, measured in pounds per square inch, reflects the difficulty of ejecting the part under mold conditions.

The Yellowness Index (Y.I.) was measured in accordance with ASTM D1825 on 5.08 cm x 7.62 cm x 0.3175 cm (2" x 3" x 1/8") chips molded at 304.4°C (580) and (650°F) 343.3°C. The magnitude of the yellowness index is an indication of the detrimental effect of an additive on the optical properties of polycarbonate. These measurements are also reported in the Tables.

The Kasha index (KI) was measured to determine the effect of the release agents on the melt stability of the resin. The KI of a resin is a measurement of its melt viscosity and is obtained in the following manner: 7 grams of resin, dried a minimum of 90 minutes at 125°C., are added to a modified Tinius-Olsen model T3 melt indexer; the temperature in the indexer is maintained at 300°C. and the resin is heated at this temperature for 6 or 12 minutes, after 6 or 12 minutes the resin is forced through a 1.0477 mm (0.04125 inch) radius orifice using a plunger of radius 4.737 mm (0.1865) inch and an applied force of 8.028 kg force of (17.7 lbs.); the time required for the plunger to travel two inches is measured in centiseconds; that is reported as the KI. For a given resin compounded with a series of different additives, the most melt stable additive (i.e. the additive which reduces the melt viscosity of the resin the least) will display the highest KI. Results are reported below in Tables 1, 2 and 3.

Other KODAR® resins can be employed in combination with the polycarbonate, for example resin prepared from terephthalic acid or ester precursor and mixtures of trans-1,4-cyclohexane dimethanol and alkylene diols having from 2 to 10 carbon atoms, preferably ethylene glycol and 1,4-butanediol. The trans-1,4-cyclohexane dimethanol is from 20 to 80 mole percent of the total diol in the polymer, the alkylene diol making up the remainder of the diol in the polymerization.

TABLE 1
BISPHENOL-A POLYCARBONATE RESIN

SAMPLE	MOLD RELEASE ¹ ADDITIVE	EJECTOR PIN PRESSURE (PSI) ²	YELLOWNESS INDEX ³	% TRANS- MISSION	KI (6 MIN)
Control 1	Pentaerythritol tetrastearate	(10,155) 64.46	---	---	---
Control 2	Eicosane (C20) ²	(3,610) 24.87	2.1	89.2	3370
Control 3	Dotricontane (C32) ²	(6,030) 41.57	1.7	90.1	3450
Control 4	Tetra tetracontane (C44) ³	(7,120) 44.95	1.7	90.1	3620
Control 5	Aldrich Mineral Oil ²	(6,250) 43.36	1.6	90.2	3480
Control 6	Drakeol 35 Mineral Oil ⁴	(9,280) 63.44	1.7	90.1	3700
Example 1	Emery 30025 (C20)	(3,975) 27.38	2.4	89.8	3280
Example 2	Emery 30045 (C31)	(4,983) 34.33	1.8	90.0	3300
Example 3	Emery 30065 (C38)	(5,836) 60.21	1.8	90.1	3280

10.3 phr mold release additive

²Purchased from Aldrich Chemical, Milwaukee, Wisconsin

³Purchased from Alfa Products, Thiabot/Ventron Division, Danvers, MA.

⁴Panreco White Mineral Oil, Pennzoil, Butler, PA.

⁵Synthetic Mineral Oil, Emery Industries, Cincinnati, Ohio

TABLE 2
BISPHENOL-A COPOLYESTERCARBONATE RESIN

<u>SAMPLE</u>	<u>MOLD RELEASE¹ ADDITIVE</u>	<u>EJECTOR PIN PRESSURE (PSI)/H₂O</u>	<u>YELLOWNESS INDEX</u>	<u>% TRANS- MISSION</u>	<u>KI (6 MIN)</u>
Control 7	none	Would not re- lease, parts broke	5.0	89.0	30945
Control 8	Pentaerythritol tetrastearate	(9380) 64.63	6.0	88.8	43930
Example 4	Emery 30025 (C20)	(6795) 46.82	4.9	89.0	45285
Example 5	Emery 30045 (C31)	(6350) 43.75	4.9	89.0	43795

¹0.3 phr mold release additive

⁵Synthetic Mineral Oil, Emery Industries, Cincinnati, Ohio

TABLE 3
80 WT. % BISPHENOL-A POLYCARBONATE/20 WT. % KODAR 150A

<u>SAMPLE</u>	<u>MOLD RELEASE¹ ADDITIVE</u>	<u>EJECTOR PIN PRESSURE (PSI) M_A</u>	<u>YELLOWNESS INDEX</u>	<u>% TRANS- MISSION</u>	<u>KI (6 MIN)</u>
Control 9	none	Would not re- lease, parts broke			
Control 10	Pentaerythritol tetrastearate	(4238) 24.20	2.7	88.3	
Example 6	Emery 30025 (C20)	(2536) 17.47	---	Visually Clear	
Example 7	Emery 30045 (C31)	(3058) 21.07	---	Visually Clear	

KI was
not
measured

10.3 phr mold release additive
 5 Synthetic Mineral Oil, Emery Industries, Cincinnati, Ohio

The ejection pressure data shows that the hydrogenated alpha-olefin oligomer fluids of the invention exhibit mold release behavior in polycarbonate, copolyesterpolycarbonate and blends of polycarbonate with alphatic polyester resin that is considerably better than the behavior of a known effective mold release agent, pentaerythritol tetrastearate. Furthermore the release properties of the hydrogenated alpha-olefin oligomer fluids in polycarbonate (Examples 1-3) are equal to or superior to the release properties of the

normal saturated alkanes of virtually equivalent molecular weight (Controls 2, 3, 4), especially when the number of carbon atoms is about 30 or greater. The data also illustrates that the mold release behavior of the hydrogenated alpha-olefin oligomer fluids is superior to that of conventional mineral oils (Controls 5 and 6). In each resin, the mold release properties exhibited by the hydrogenated alpha-olefin oligomer fluids is achieved without experiencing a loss in optical properties or melt viscosity.

Claims

1. A composition comprising an aromatic carbonate polymer in admixture with a mold release effective amount of a hydrogenated alpha-olefin oligomer fluid.
2. The composition in accordance with claim 1 wherein the aromatic carbonate polymer is a polycarbonate.
3. The composition in accordance with claim 2 wherein the polycarbonate is bisphenol-A polycarbonate.
4. The composition in accordance with claim 1 wherein the aromatic carbonate polymer is a copolyester-carbonate.
5. The composition in accordance with claim 1 wherein 0.01 to 1.0 weight percent of the hydrogenated alpha-olefin oligomer fluid is present.
6. The composition in accordance with claim 5 wherein 0.05 to 0.5 weight percent of the hydrogenated alpha-olefin oligomer fluid is present.
7. The composition in accordance with claim 1 wherein the hydrogenated alpha-olefin oligomer fluid is prepared from normal alpha-olefins having from six to twelve carbon atoms, inclusive.
8. The composition in accordance with claim 1 wherein the normal alpha-olefin is decene-1.
9. The composition in accordance with claim 1 wherein the hydrogenated alpha-olefin oligomer fluid has at least 30 carbon atoms.
10. The composition in accordance with claim 1 wherein also present in the composition up to 50 weight percent is a polyester derived from terephthalic acid or ester precursor or a mixture of terephthalic acid or ester precursor or a mixture of terephthalic and isophthalic acid or precursors and trans-1,4-cyclohexane dimethanol or a mixture of trans-1,4-cyclohexane dimethanol and an alkylene diol of 2 to 10 carbon atoms, inclusive.

Revendications

1. Composition comprenant un polymère de carbonate aromatique en mélange avec une quantité efficace pour le démoulage d'un fluide d'oligomère d'alpha-oléfine hydrogéné.
2. Composition selon la revendication 1, dans laquelle le polymère de carbonate aromatique est un polycarbonate.
3. Composition selon la revendication 2, dans laquelle le polycarbonate est du polycarbonate de bisphénol-A.
4. Composition selon la revendication 1, dans laquelle le polymère de carbonate aromatique est un copolyestercarbonate.
5. Composition selon la revendication 1, dans laquelle de 0,01 à 1,0 pourcent en poids du fluide d'oligomère d'alpha-oléfine hydrogéné est présent.
6. Composition selon la revendication 5, dans laquelle de 0,05 à 0,5 pourcent en poids du fluide d'oligomère d'alpha-oléfine hydrogéné est présent.

7. Composition selon la revendication 1, dans laquelle on prépare le fluide d'oligomère d'alpha-oléfine hydrogéné à partir d'alpha-oléfines normales ayant de six à douze atomes de carbone, inclus.
8. Composition selon la revendication 1, dans laquelle l'alpha-oléfine normale est le décène-1.
- 5 9. Composition selon la revendication 1, dans laquelle le fluide d'oligomère d'alpha-oléfine hydrogéné contient au moins 30 atomes de carbone.
- 10 10. Composition selon la revendication 1, dans laquelle est également présent dans la composition jusqu'à 50 pourcent en poids d'un polyester dérivé d'acide téréphtalique ou de précurseur d'ester ou d'un mélange d'acide téréphtalique ou de précurseur d'ester ou d'un mélange d'acides téréphtalique et isophtalique ou de précurseurs et de trans-dihydroxyméthyl-1,4 cyclohexane ou d'un mélange de trans-dihydroxyméthyl-1,4 cyclohexane et d'un alkylènediol ayant de 2 à 10 atomes de carbone, inclus.

15 Patentansprüche

1. Zusammensetzung enthaltend ein aromatisches Carbonatpolymer in Mischung mit einer wirksamen Menge eines Formentrennmittels aus einem hydrierten Alpha-Olefin Oligomerfluid.
- 20 2. Zusammensetzung nach Anspruch 1, worin das aromatische Carbonatpolymer ein Polycarbonat ist.
3. Zusammensetzung nach Anspruch 2, worin das Polycarbonat Bisphenol-A Polycarbonat ist.
4. Zusammensetzung nach Anspruch 1, worin das aromatische Carbonatpolymer ein Copolyestercarbonat ist.
- 25 5. Zusammensetzung nach Anspruch 1, worin 0,01 - 1,0 Gew.-% des hydrierten Alpha-Olefin Oligomerfluids vorhanden sind.
- 30 6. Zusammensetzung nach Anspruch 5, worin 0,05 bis 0,5 Gew.-% des hydrierten Alpha-Olefin Oligomerfluids vorhanden sind.
7. Zusammensetzung nach Anspruch 1, worin das hydrierte Alpha-Olefin Oligomerfluid aus normalen Alpha-Olefinen mit 6 bis 12 Kohlenstoffatomen einschließlich hergestellt ist.
- 35 8. Zusammensetzung nach Anspruch 1, worin das normale Alpha-Olefin Decen-1 ist.
9. Zusammensetzung nach Anspruch 1, worin das hydrierte Alpha-Olefin Oligomerfluid wenigstens 30 Kohlenstoffatome aufweist.
- 40 10. Zusammensetzung nach Anspruch 1, worin ebenfalls enthalten sind bis zu 50 Gew.-% eines Polyesters, der von Terephthalsäure oder dem Estervorläufer oder einer Mischung von Terephthalsäure oder Estervorläufer oder einer Mischung von Terephthalsäure und Isophthalsäure oder Vorläufern und Trans-1,4-cyclohexandimethanol oder einer Mischung von Trans-1,4-cyclohexandimethanol und einem Alky-
45 lëndiol mit 2 bis 10 Kohlenstoffatomen einschließlich abgeleitet ist.

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